In the claims:

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- 1. (currently amended) A plastid transformation and expression vector useful for transformation of a target plastid of a higher plant species, said vector comprising an expression cassette which comprises an expression cassette comprising as operably linked components, a 5" part of the plastid DNA sequence inclusive of the spacer sequence, a promoter operative in said plastids the target plastid, a selectable marker sequence, at least one DNA sequence encoding at least [a]an immunologically active portion of an immunoglobulin mutimeric multimeric chain, at least one DNA sequence encoding a chaperonin, a transcription termination region functional in said the target plastid, and the 3'on each of the 5' and 3' ends of said expression cassette DNA sequences which are homologous to a part of the target plastid DNA sequence genome of a higher plant species.
- 2. (currently amended) The plastid transformation and expression vector of claim 1, wherein said immunoglobulin mutimeric multimeric chain comprises a heavy chain.
- 3. (currently amended) The plastid transformation and expression vector of claim 1, wherein said immunoglobulin mutimeric multimeric chain comprises a light chain.
- 4. (currently amended) The plastid transformation and expression vector of claim 1, wherein said immunoglobulin mutimeric multimeric chain comprises both a heavy and a light chain.
- 5. (currently amended) The plastid transformation and expression vector of claim 1, wherein said immunoglobulin mutimeric multimeric chain comprises a single-chain variable fragment (scFv).
- 6. (currently amended) The plastid transformation and expression vector of claim 1, wherein said immunoglobulin mutimeric multimeric chain comprises a heavy chain constant region fused to an operative ligand.
- 7. (currently amended) The plastid transformation and expression vector of claim 4, wherein said heavy and light chains are separated by a linker comprising an intervening a stop codon and a ribosome binding site.

Claims 8-27 (canceled)

28. (currently amended) A method for introducing DNA encodingproducing immunoglobulin mutimeric chain coding sequences intoprotein in a plastid, said method comprising:

introducing a plastid expression vector into a plant cell[,] of a higher plant species having a target plastid expression vector adsorbed onto a microprojectile,

said plastid expression vector comprising as operably linked components[,]

a DNA sequence containing at least one plastid replication origin functional in [a]the target plastid,

a transcriptional initiation region functional in [a]the target plastid,

at least one heterologous DNA sequence encoding at least [a]an immunologically active portion of an immunoglobulin mutimeric multimeric chain,

at least one DNA sequence encoding a chaperonin and

a transcriptional termination region functional in said cellsthe target plasmid, whereby said heterologous DNA is introduced into [a]the target plastid in saidthe plant cell; and wherein a multimeric immunoglobulin is produced.

- 29. (currently amended) The method of claim 28, wherein said immunoglobulin mutimeric multimeric chain comprises a heavy chain.
- 30. (currently amended) The method of claim 28, wherein said immunoglobulin mutimeric multimeric chain comprises a light chain.
- 31. (currently amended) The method of claim 28, wherein said immunoglobulin mutimeric multimeric chain comprises both a heavy chain and a light chain.
- 32. (currently amended) The method of claim 28, wherein said immunoglobulin mutimeric multimeric chain comprises a single-chain variable fragment (scFv).

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33. (currently amended) The method of claim 28, wherein said immunoglobulin mutimeric multimeric chain comprises a heavy chain constant region fused to an operative ligand.

Claims 34-50 (canceled)